



## Complete Summary

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### GUIDELINE TITLE

Stress related conditions and other mental disorders.

### BIBLIOGRAPHIC SOURCE(S)

Work Loss Data Institute. Stress related conditions and other mental disorders. Corpus Christi (TX): Work Loss Data Institute; 2005. 104 p. [123 references]

### GUIDELINE STATUS

Note: This guideline has been updated. The National Guideline Clearinghouse (NGC) is working to update this summary.

### \*\* REGULATORY ALERT \*\*

### FDA WARNING/REGULATORY ALERT

Note from the National Guideline Clearinghouse: This guideline references a drug(s) for which important revised regulatory and/or warning information has been released.

- On October 17, 2005, Eli Lilly and the U.S. Food and Drug Administration (FDA) notified healthcare professionals of revision to the PRECAUTIONS/Hepatotoxicity section of the prescribing information for Cymbalta (duloxetine hydrochloride), indicated for treatment of major depressive disorder and diabetic peripheral neuropathic pain. Postmarketing reports of hepatic injury (including hepatitis and cholestatic jaundice) suggest that patients with preexisting liver disease who take duloxetine may have an increased risk for further liver damage. The new labeling extends the Precaution against using Cymbalta in patients with substantial alcohol use to include those patients with chronic liver disease. It is recommended that Cymbalta not be administered to patients with any hepatic insufficiency. See the [FDA Web site](#) for more information.
- On July 1, 2005, in response to recent scientific publications that report the possibility of increased risk of suicidal behavior in adults treated with antidepressants, the U.S. Food and Drug Administration (FDA) issued a Public Health Advisory to update patients and healthcare providers with the latest information on this subject. Even before the publication of these recent reports, FDA had already begun the process of reviewing available data to determine whether there is an increased risk of suicidal behavior in adults taking antidepressants. The Agency has asked manufacturers to provide information from their trials using an approach similar to that used in the evaluation of the risk of suicidal behavior in the pediatric population taking antidepressants. This effort will involve hundreds of clinical trials and may

take more than a year to complete. See the [FDA Web site](#) for more information.

## COMPLETE SUMMARY CONTENT

\*\* REGULATORY ALERT \*\*

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## SCOPE

### DISEASE/CONDITION(S)

Work-related stress and other mental disorders

### GUIDELINE CATEGORY

Counseling

Diagnosis

Evaluation

Management

Treatment

### CLINICAL SPECIALTY

Family Practice

Internal Medicine

Psychiatry

Psychology

### INTENDED USERS

Advanced Practice Nurses

Health Care Providers

Health Plans

Nurses

Physician Assistants

Physicians

Psychologists/Non-physician Behavioral Health Clinicians

### GUIDELINE OBJECTIVE(S)

To offer evidence-based step-by-step decision protocols for the assessment and treatment of workers' compensation conditions

## TARGET POPULATION

Workers with occupational stress and other mental disorders

## INTERVENTIONS AND PRACTICES CONSIDERED

The following interventions were considered and recommended as indicated in the original guideline document:

1. Acceptance and commitment therapy (ACT)
2. Activity restrictions/Work modifications
3. Acupressure
4. Antidepressants (the choice of first-line therapy between selective serotonin reuptake inhibitors [SSRIs] and tricyclic antidepressants [TCA] is currently under study)
5. Aromatherapy
6. Cognitive therapy
7. Cognitive behavioral stress management (CBSM)
8. Depression screening
9. Disease management programs
10. Distractive methods
11. Duloxetine (Cymbalta®)
12. Patient education
13. Electroconvulsive therapy (ECT)
14. Exercise
15. Kava extract
16. Light therapy
17. Massage therapy
18. Mind/body interventions (relaxation)
19. Music
20. Peer support
21. Psychosocial empowerment programs
22. Return to work
23. St. John's wort
24. Stress inoculation training
25. Stress management, behavioral/cognitive interventions
26. Stress management, physical interventions (aerobic exercise)
27. Therapist optimism
28. Vitamin use (multi-vitamin and mineral supplements)
29. Yoga

The following interventions/procedures are under study and are not specifically recommended:

1. Acupuncture
2. Brain wave synchronizers for stress reduction
3. Computer-assisted cognitive therapy
4. Expatriate employee adjustment
5. Fatigue (as precursor to stress)

6. Folate
7. Hypnosis
8. Innovative promotion program
9. Opioid antagonists (naltrexone)
10. Pharmaceuticals versus behavioral therapy for tension headaches
11. Psychosocial and pharmacological treatment (for deliberate self harm)
12. Technological stress

The following interventions/procedures were considered, but are not recommended:

1. Psychological debriefing (for preventing post-traumatic stress disorder)
2. Vitamin B6

## MAJOR OUTCOMES CONSIDERED

Effectiveness of treatments in reducing stress and anxiety

## METHODOLOGY

### METHODS USED TO COLLECT/SELECT EVIDENCE

Hand-searches of Published Literature (Primary Sources)  
Searches of Electronic Databases

### DESCRIPTION OF METHODS USED TO COLLECT/SELECT THE EVIDENCE

Not stated

### NUMBER OF SOURCE DOCUMENTS

Not stated

### METHODS USED TO ASSESS THE QUALITY AND STRENGTH OF THE EVIDENCE

Weighting According to a Rating Scheme (Scheme Given)

### RATING SCHEME FOR THE STRENGTH OF THE EVIDENCE

Ranking by quality within type of evidence:

- a. High Quality
- b. Medium Quality
- c. Low Quality

### METHODS USED TO ANALYZE THE EVIDENCE

Review of Published Meta-Analyses  
Systematic Review

## DESCRIPTION OF THE METHODS USED TO ANALYZE THE EVIDENCE

Not stated

## METHODS USED TO FORMULATE THE RECOMMENDATIONS

Not stated

## RATING SCHEME FOR THE STRENGTH OF THE RECOMMENDATIONS

Not applicable

## COST ANALYSIS

A formal cost analysis was not performed and published cost analyses were not reviewed.

## METHOD OF GUIDELINE VALIDATION

Not stated

## DESCRIPTION OF METHOD OF GUIDELINE VALIDATION

Not applicable

## RECOMMENDATIONS

### MAJOR RECOMMENDATIONS

Note: This guideline has been updated. The National Guideline Clearinghouse (NGC) is working to update this summary. The recommendations that follow are based on the previous version of the guideline.

#### Initial Diagnosis

Stress is the most common mental condition treated by occupational or primary care physicians and will be the focus of this guideline. References to additional mental disorders are found in the procedure summary in the original guideline document, although the more severe of those usually require referral to a specialist. Stress is not its own diagnosis but rather a combination of nonspecific emotional or physical symptoms varying in intensity and duration, which may or may not be related to a specific incident. The stress might also be associated with a particular disease or syndrome, but that is not always the case.

A stressor is defined as anything that exerts a physical, emotional, or mental demand on an individual. Stress often occurs when the individual has anxiety because of a mismatch between perceived demands and resources, whether work-related or personal. The source of stress can be acute (such as an employee relocation) or chronic (such as consistently poor relations with a supervisor).

For some people, stress causes or contributes to a deterioration of physical health, resulting in more headaches or more common colds. While the scientific literature is not clear on this topic, stress may also contribute to the worsening of more serious conditions such as heart disease, irritable bowel disease, or ulcers, depending on the individual's coping methods. On the other hand, the presence of certain physical conditions could be the cause of stress.

## Initial Evaluation

Focus on identifying possible red flags or warning signs for potentially serious psychopathology that would require immediate specialty referral. Red flags may include impairment of mental functions, overwhelming symptoms, signs of substance abuse, or debilitating depression. In the absence of red flags, the occupational or primary care physician can handle most common stress-related conditions safely.

In talking to the patient, it is important for the physician to get him or her to try and explain or pinpoint incidents or reasons for the stress, rather than to just generalize (i.e., "I hate my job," "Everything makes me stressed out"). The physician may have to ask more specific questions about work or home life if the patient is initially unwilling or unable to address specific issues.

Occupational stress usually stems from one of three common models:

1. Person-environment fit model: Poor job fit, such as a mismatch between the skills of the individual and the demands of the job, or a disparity between the individual's career-related desires versus actual opportunities presented, is a leading cause of workplace stress.
2. Demand control model: Jobs that place high demands on the worker but give him or her little control or opportunities for decision-making lead to high job strain, a source of stress that is consistently linked as a contributor to physical conditions such as cardiovascular mortality, heart disease, and hypertension. Consideration should be given to the influence of the individual's occupational and personal history, which may have an effect on how this model applies to his or her situation.
3. Effort-reward model: Shows that stress is often the result of high effort without social reward. Like the demand control model, this model points out that a low ratio of effort to reward leads to sustained autonomic arousal and can cause physical effects such as high blood pressure or myocardial infarction.

Exploration of how and if the patient's stress follows the path of one of the above models will be helpful in determining treatment.

More specific sources of stress include bereavement, illness, familial changes or disorder, or other common and/or traumatic life changes. Time off work may be helpful, although the ultimate goal should be to preserve the patient's ability to function both occupationally and socially. Time off should not be so excessive that the employee loses his or her sense of function and appreciation at work and at home.

## Initial Therapy

1. Pursuing the patient's thoughts on how his or her stress relates to the above models may help determine the source of stress and cultivate ideas on how to eliminate or cope with the stress. Patient education and understanding about stress is necessary for effective stress management to take place.
2. Other common treatment pathways include the use of one or more of the following:
  - a. Relaxation techniques (such as meditation)
  - b. Exercise (aerobic exercise has been shown to positively influence mood)
  - c. Behavioral training (such as time management, anger management, assertiveness, or conflict resolution training)
  - d. Stress inoculation therapy
  - e. Cognitive therapy
  - f. Modified work
  - g. Organizational interventions
3. Pharmaceutical therapy (limited, short-term use of anti-anxiety agents to improve function--anything else should be used in conjunction with a specialty referral)

Follow-up visits are an important part of treatment and should be conducted by a mid-level practitioner in person or via phone every three or four days, depending on the severity of the case, while a path to recognizable treatment is established and followed. Failure to improve or make significant progress after several months may indicate the need for psychiatric assessment or counseling.

#### Official Disability Guidelines (ODG) Return-To-Work Pathways

Senile and Presenile Organic Psychotic Conditions (see original guideline document for ICD-9 codes for this and other diagnoses)

Not severe, medical treatment: 0 days

Severe, specially designed, limited modified work: 7 days

Severe, regular work: indefinite

#### Senile Dementia with Delusional or Depressive Features

Severe, specially designed, limited modified work: 7 days

Severe, affecting fellow worker productivity & safety: indefinite

Severe, regular work: indefinite

#### Alcohol Withdrawal Delirium

Without hospitalization: 1-7 days

Including rehab, substance abuse professional (SAP) evaluation: 28 days

Including rehab, SAP evaluation, job safety issues: 42 days

#### Drug Withdrawal Syndrome

Without hospitalization: 0-5 days

With hospitalization, without suicidal ideation: 7 days

With hospitalization, with suicidal ideation: 21 days

#### Paranoid and/or Hallucinatory States Induced by Drugs

Without hospitalization: 1-3 days

With hospitalization, without threat of harm: 7 days

With hospitalization, with threat of harm: 21 days

#### Transient Organic Psychotic Conditions

14 days

#### Paranoid Type

Without hospitalization, no job safety issues: 0-7 days

With hospitalization: 42 days or by report

#### Unspecified Schizophrenia

Without hospitalization, no job safety issues: 0-7 days

With hospitalization: 16-42 days

#### Major Depressive Disorder, Single Episode

Rule out impaired mood/personality disorder: 0 days

Outpatient therapy, without symptoms affecting work: 0-7 days

Outpatient therapy, with symptoms interfering with work: 21-42 days

With hospitalization, non-cognitive/modified work: 21 days

With hospitalization, cognitive work: 42 days

#### Major Depressive Disorder, Recurrent Episode



Outpatient therapy, without symptoms affecting work: 0-7 days

Outpatient therapy, with symptoms interfering with work: 14-28 days

With hospitalization, non-cognitive/modified work: 21 days

With hospitalization, cognitive work: 42 days

#### Bipolar Affective Disorder, Depressed

Rule out impaired mood/personality disorder: 0 days

Without hospitalization: 0-21 days

With hospitalization: 21-42 days

#### Bipolar Affective Disorder, Mixed

Without hospitalization: 0-14 days

With hospitalization: 21-42 days

#### Paranoia

Without hospitalization: 0-14 days

With hospitalization: 14-21 days

#### Depressive Type Psychosis

Without hospitalization: 0-56 days

With hospitalization: 21-64 days

#### Anxiety States

Rule out impaired mood/personality disorder: 0 days

Without hospitalization: 0-7 days

With hospitalization: 14-21 days

#### Panic Disorder

1-14 days

#### Generalized Anxiety Disorder

14-21 days

#### Hysteria

Without hospitalization: 0 days

With hospitalization: 7-14 days

#### Obsessive-Compulsive Disorders

Without hospitalization: 0 days

With hospitalization: 10 days

#### Personality Disorders

0 days

#### Alcohol Dependence Syndrome

Without hospitalization: 1 day

Without hospitalization, considering fellow worker danger & morale: 7-14 days

With hospitalization, including rehab: 14-28 days

Safety sensitive position: as determined by the SAP

#### Acute Alcoholic Intoxication

1-2 days

Also treated as rule violation absence

#### Opioid Type Dependence

Without hospitalization: 0 days

Without hospitalization, considering fellow worker danger & morale: 7-14 days

With hospitalization, including rehab: 14-38 days (10 days post-discharge)

Safety sensitive position: as determined by the SAP

#### Barbiturate and Similarly Acting Sedative or Hypnotic Dependence

Without hospitalization: 0 days

Without hospitalization, considering fellow worker danger & morale: 7-14 days

With hospitalization: 21 days

With hospitalization, plus rehab: 28 days

Safety sensitive position: as determined by the SAP

#### Cocaine Dependence

Without hospitalization: 0 days

Without hospitalization, considering fellow worker danger & morale: 7-14 days

With hospitalization: 28 days

Safety sensitive position: as determined by the SAP

#### Cannabis Dependence

0-2 days

#### Amphetamine and Other Psychostimulant Dependence

Without hospitalization: 0 days

Without hospitalization, considering fellow worker danger & morale: 7-14 days

With hospitalization: 14 days

With hospitalization, plus rehab: 28 days

Safety sensitive position: as determined by the SAP

#### Hallucinogen Dependence

Without hospitalization: 0 days

Without hospitalization, considering fellow worker danger & morale: 7-14 days

With hospitalization: 10 days

With hospitalization, plus rehab: 28 days

Safety sensitive position: as determined by the SAP

#### Alcohol Abuse

1 day

#### Cocaine Abuse

Without hospitalization: 0-1 days

With hospitalization: 10 days

With hospitalization, plus rehab: 28 days

#### Amphetamine or Related Acting Sympathomimetic Abuse

Without hospitalization: 1 day

With hospitalization: 14 days

With hospitalization, plus rehab: 28 days

#### Acute Reaction to Stress

Without hospitalization (on-going counseling/drug therapy): 1 day

With hospitalization: 10 days

#### Unspecified Acute Reaction to Stress, Post-traumatic Stress Disorder

Without hospitalization (on-going counseling): 1 day

With hospitalization: 10 days

Chemical dependence comorbidity: 28 days

#### Adjustment Reaction

Without hospitalization: 1-6 days

Outpatient care: 1-6 days

With inpatient hospitalization: 14-28 days

#### Postconcussion Syndrome

Mild: 1 day

Severe: by report

#### Depressive Disorder, not Elsewhere Classified

Rule out impaired mood/personality disorder: 0 days

Outpatient therapy, without symptoms affecting work or other job issues: 0-7 days

Outpatient therapy, with symptoms interfering with work: 21 days

Outpatient therapy, with serious job satisfaction issues: 28-42 days

With hospitalization, non-cognitive/modified work: 28 days

With hospitalization, cognitive work: 42-56 days

#### Attention Deficit Disorder

1 day

(See ODG Capabilities & Activity Modifications for Restricted Work under "Work" in the Procedure Summary of the original guideline document)

#### CLINICAL ALGORITHM(S)

None provided

### EVIDENCE SUPPORTING THE RECOMMENDATIONS

#### TYPE OF EVIDENCE SUPPORTING THE RECOMMENDATIONS

During the comprehensive medical literature review, preference was given to high quality systematic reviews, meta-analyses, and clinical trials over the past ten years, plus existing nationally recognized treatment guidelines from the leading specialty societies.

The type of evidence associated with each recommended or considered intervention or procedure is ranked in the guideline's annotated reference summaries.

Ranking by Type of Evidence:

1. Systematic Review/Meta-Analysis
2. Controlled Trial-Randomized (RCT) or Controlled
3. Cohort Study-Prospective or Retrospective
4. Case Control Series
5. Unstructured Review
6. Nationally Recognized Treatment Guideline (from [www.guideline.gov](http://www.guideline.gov))
7. State Treatment Guideline
8. Foreign Treatment Guideline
9. Textbook
10. Conference Proceedings/Presentation Slides

## BENEFITS/HARMS OF IMPLEMENTING THE GUIDELINE RECOMMENDATIONS

### POTENTIAL BENEFITS

These guidelines unite evidence-based protocols for medical treatment with normative expectations for disability duration. They also bridge the interests of the many professional groups involved in diagnosing and treating work-related stress and other mental disorders.

### POTENTIAL HARMS

Not stated

## IMPLEMENTATION OF THE GUIDELINE

### DESCRIPTION OF IMPLEMENTATION STRATEGY

An implementation strategy was not provided.

## INSTITUTE OF MEDICINE (IOM) NATIONAL HEALTHCARE QUALITY REPORT CATEGORIES

### IOM CARE NEED

Getting Better

### IOM DOMAIN

Effectiveness

## IDENTIFYING INFORMATION AND AVAILABILITY

### BIBLIOGRAPHIC SOURCE(S)

Work Loss Data Institute. Stress related conditions and other mental disorders. Corpus Christi (TX): Work Loss Data Institute; 2005. 104 p. [123 references]

### ADAPTATION

Not applicable: The guideline was not adapted from another source.

### DATE RELEASED

2004 (revised 2005)

### GUIDELINE DEVELOPER(S)

Work Loss Data Institute - Public For Profit Organization

#### SOURCE(S) OF FUNDING

Not stated

#### GUIDELINE COMMITTEE

Not stated

#### COMPOSITION OF GROUP THAT AUTHORED THE GUIDELINE

Not stated

#### FINANCIAL DISCLOSURES/CONFLICTS OF INTEREST

Not stated

#### GUIDELINE STATUS

Note: This guideline has been updated. The National Guideline Clearinghouse (NGC) is working to update this summary.

#### GUIDELINE AVAILABILITY

Electronic copies of the updated guideline: Available to subscribers from the [Work Loss Data Institute Web site](#).

Print copies: Available from the Work Loss Data Institute, 169 Saxony Road, Suite 210, Encinitas, CA 92024; Phone: 800-488-5548, 760-753-9992, Fax: 760-753-9995; [www.worklossdata.com](http://www.worklossdata.com).

#### AVAILABILITY OF COMPANION DOCUMENTS

Background information on the development of the Official Disability Guidelines of the Work Loss Data Institute is available from the [Work Loss Data Institute Web site](#).

#### PATIENT RESOURCES

None available

#### NGC STATUS

This NGC summary was completed by ECRI on April 4, 2005. This summary was updated by ECRI on August 15, 2005, following the U.S. Food and Drug Administration advisory on antidepressant medications. This summary was updated by ECRI on October 20, 2005, following the U.S. Food and Drug Administration advisory on Cymbalta (duloxetine hydrochloride). This NGC summary was updated by ECRI on January 30, 2006.

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